Louisiana Medicaid
Osteoporosis – Bone Resorption Suppression Agents

The *Louisiana Uniform Prescription Drug Prior Authorization Form* should be utilized to request prior authorization for non-preferred bone resorption suppression agents.

Additional Point-of-Sale edits may apply.

*These agents may have **Black Box Warnings** and/or may be subject to **Risk Evaluation and Mitigation Strategy (REMS)** under FDA safety regulations. Please refer to individual prescribing information for details.*

Approval Criteria for Initial and Reauthorization Requests

- There is no preferred alternative that is the exact same chemical entity, formulation, strength, etc.; **AND**
- Previous use of a preferred product - **ONE** of the following is required:
    - The recipient has had a *treatment failure* with at least one preferred product; **OR**
    - The recipient has had an *intolerable side effect* to at least one preferred product; **OR**
    - The recipient has *documented contraindication(s)* to the preferred products that are appropriate to use for the condition being treated; **OR**
    - There is *no preferred product that is appropriate* to use for the condition being treated; **OR**
    - The prescriber states that the recipient is currently using the requested medication; **AND**
- If the request is for abaloparatide (Tymlos®) or teriparatide (Forteo®);
    - Begin and end dates of previous treatment for abaloparatide and/or teriparatide are **stated on the request**; **AND**
    - The recipient has not used either of these agents for a cumulative duration of 24 months during the recipient's lifetime; **AND**
- By submitting the authorization request, the prescriber attests to the following:
    - The prescribing information for the requested medication has been thoroughly reviewed, including any Black Box Warning, Risk Evaluation and Mitigation Strategy (REMS), contraindications, minimum age requirements, recommended dosing, and prior treatment requirements; **AND**
    - All laboratory testing and clinical monitoring recommended in the prescribing information have been completed as of the date of the request and will be repeated as recommended; **AND**
    - The recipient has no concomitant drug therapies or disease states that limit the use of the requested medication and will not be receiving the requested medication in combination with any other medication that is contraindicated or not recommended per FDA labeling.

Duration of initial and reauthorization approval: 12 months

References

Clinical Pharmacology [database online]. Tampa, FL: Gold Standard, Inc.; Retrieved from https://www.clinicalkey.com/pharmacology/

DiPiro JT, Talbert RL, Yee GC, Matzke GR, Wells BG, Posey L. eds. Pharmacotherapy: A Pathophysiologic Approach, 10e New York, NY: McGraw-Hill; Retrieved from https://accesspharmacy.mhmedical.com/book.aspx?bookid=1861

Forteo (teriparatide) [package insert]. Indianapolis, IN: Eli Lilly and Company; November 2020. http://uspl.lilly.com/forteo/forteo.html#pi

Tymlos (abaloparatide) [package insert]. Waltham, MA: Radius Health, Inc; October 2020. http://radiuspharm.com/wp-content/uploads/tymlos/tymlos-prescribing-information.pdf

Revision / Date	**Implementation Date**
Single PDL Implementation	May 2019
Separated "Select Therapeutic Classes (Established)" into individual therapeutic class documents / November 2019	January 2020
Formatting changes, updated references / April 2021	July 2021